

<b>NCT ID</b>	NCT03084471
<b>Title</b>	An Open-Label, Multi-Centre, Study to Assess the Safety of Fixed-Dose Durvalumab + Tremelimumab Combination Therapy or Durvalumab Monotherapy in Advanced Solid Malignancies.
<b>Phase</b>	Phase 3
<b>Date Added</b>	2017-03-21
<b>Location</b>	California, United States District of Columbia, United States Illinois, United States Nebraska, United States New Jersey, United States New York, United States South Carolina, United States Tennessee, United States Virginia, United States Washington, United States Canada France Germany Italy Korea, Republic of Netherlands United Kingdom
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	No
<b>CRC-directed</b>	No
<b>Status</b>	Active, not recruiting
<b>Drugs</b>	MEDI4736 (Durvalumab), MEDI4736 (Durvalumab) + Tremelimumab

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<b>NCT ID</b>	NCT05198934
<b>Title</b>	Sotorasib and Panitumumab Versus Investigator's Choice for Participants With Kirsten Rat Sarcoma (KRAS) p.G12C Mutation (CodeBreak 300)
<b>Phase</b>	Phase 3
<b>Date Added</b>	2022-01-20
<b>Location</b>	Alabama, United States California, United States District of Columbia, United States Florida, United States Georgia, United States Michigan, United States New York, United States North Carolina, United States Ohio, United States Pennsylvania, United States Tennessee, United States Texas, United States Australia France Germany Greece Italy Japan Korea, Republic of Mexico Spain Taiwan United Kingdom
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	Yes
<b>CRC-directed</b>	Yes
<b>Status</b>	Recruiting
<b>Drugs</b>	

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<b>NCT ID</b>	NCT05141721
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**Title** A Study of a Personalized Neoantigen Vaccine in Combination With Immune Checkpoint Blockade for Patients With Metastatic Colorectal Cancer

**Phase** Phase 2, Phase 3

**Date Added** 2021-12-02

**Location** Arizona, United States  
Arkansas, United States  
California, United States  
Colorado, United States  
Connecticut, United States  
Florida, United States  
Illinois, United States  
Indiana, United States  
Kansas, United States  
Kentucky, United States  
Maryland, United States  
Michigan, United States  
Nevada, United States  
New Jersey, United States  
New York, United States  
North Carolina, United States  
Ohio, United States  
Oregon, United States  
Pennsylvania, United States  
South Carolina, United States  
Tennessee, United States  
Texas, United States  
Utah, United States  
Virginia, United States  
Washington, United States  
Wisconsin, United States

**IO -centered** No

**Prior IO Allowed** No

**CRC-directed** Yes

**Status** Recruiting

**Drugs** Atezolizumab, Bevacizumab, Fluoropyrimidine, GRT-C901, GRT-R902, Ipilimumab, oxaliplatin

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**NCT ID** NCT04607421

**Title** A Study of Encorafenib Plus Cetuximab With or Without Chemotherapy in People With Previously Untreated Metastatic Colorectal Cancer (BREAKWATER)

**Phase** Phase 3

**Date Added** 2020-10-29

<b>Location</b>	Arizona, United States California, United States District of Columbia, United States Florida, United States Georgia, United States Illinois, United States Louisiana, United States Michigan, United States Minnesota, United States Missouri, United States Montana, United States Nebraska, United States New Jersey, United States New York, United States North Carolina, United States Ohio, United States Oklahoma, United States Oregon, United States Pennsylvania, United States Tennessee, United States Texas, United States Virginia, United States Washington, United States Wisconsin, United States Argentina Australia Belgium Brazil Bulgaria Canada China Czechia Denmark Finland Germany India Italy Japan Korea, Republic of Mexico Netherlands New Zealand Norway Poland Russian Federation Slovakia South Africa Spain Sweden Taiwan Ukraine United Kingdom
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	No
<b>CRC-directed</b>	Yes
<b>Status</b>	Recruiting
<b>Drugs</b>	5-FU, Bevacizumab, capecitabine, cetuximab, encorafenib, Irinotecan, Leucovorin, oxaliplatin
<b>NCT ID</b>	NCT05328908
<b>Title</b>	A Study of Nivolumab-relatlimab Fixed-dose Combination Versus Regorafenib or TAS-102 in Participants With Later-lines of Metastatic Colorectal Cancer
<b>Phase</b>	Phase 3
<b>Date Added</b>	2022-04-14

<b>Location</b>	Arkansas, United States California, United States Connecticut, United States Florida, United States Georgia, United States Idaho, United States Illinois, United States Indiana, United States Massachusetts, United States Michigan, United States Minnesota, United States New Jersey, United States North Carolina, United States Ohio, United States Pennsylvania, United States South Carolina, United States South Dakota, United States Tennessee, United States Texas, United States Virginia, United States Wisconsin, United States Argentina Australia Austria Belgium Canada Chile China Czechia France Germany Italy Japan Korea, Republic of Netherlands Poland Puerto Rico Singapore Spain Sweden Switzerland Taiwan
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	No
<b>CRC-directed</b>	Yes
<b>Status</b>	Recruiting
<b>Drugs</b>	Nivolumab-relatlimab FDC, Regorafenib, TAS-102, Lonsurf, Opdivo, Stivarga
<b>NCT ID</b>	NCT05223673
<b>Title</b>	Phase 3 Study of Futuximab/Modotuximab in Combination With Trifluridine/Tipiracil Versus Trifluridine/Tipiracil Single Agent in Participants With Previously Treated Metastatic Colorectal Cancer
<b>Phase</b>	Phase 3
<b>Date Added</b>	2022-02-04
<b>Location</b>	Michigan, United States Ohio, United States Belgium Denmark Finland Japan
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	No
<b>CRC-directed</b>	Yes
<b>Status</b>	Recruiting
<b>Drugs</b>	Futuximab/modotuximab, Trifluridine/Tipiracil

<b>NCT ID</b>	NCT02928224
<b>Title</b>	Study of Encorafenib + Cetuximab Plus or Minus Binimetinib vs. Irinotecan/Cetuximab or Infusional 5-Fluorouracil (5-FU)/Folinic Acid (FA)/Irinotecan (FOLFIRI)/Cetuximab With a Safety Lead-in of Encorafenib + Binimetinib + Cetuximab in Patients With BRAF V600E-mutant Metastatic Colorectal Cancer
<b>Phase</b>	Phase 3
<b>Date Added</b>	2016-10-10
<b>Location</b>	Arizona, United States California, United States Colorado, United States Connecticut, United States Florida, United States Illinois, United States Indiana, United States Iowa, United States Kansas, United States Louisiana, United States Maryland, United States Massachusetts, United States Minnesota, United States Missouri, United States New Jersey, United States New York, United States Ohio, United States Oregon, United States Tennessee, United States Texas, United States Washington, United States Wisconsin, United States Argentina Australia Austria Belgium Brazil Canada Chile Czechia Denmark France Germany Hungary Israel Italy Japan Korea, Republic of Mexico Netherlands Norway Poland Russian Federation Spain Taiwan Turkey Ukraine United Kingdom
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	Yes
<b>CRC-directed</b>	Yes
<b>Status</b>	Active, not recruiting
<b>Drugs</b>	5-Fluorouracil, Binimetinib, cetuximab, encorafenib, Folinic acid, Irinotecan

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<b>NCT ID</b>	NCT04607668
<b>Title</b>	Trilaciclib, a CDK 4/6 Inhibitor, in Patients Receiving FOLFOXIRI/Bevacizumab for Metastatic Colorectal Cancer (mCRC):
<b>Phase</b>	Phase 3
<b>Date Added</b>	2020-10-29

**Location**  
Arizona, United States  
California, United States  
District of Columbia, United States  
Florida, United States  
Georgia, United States  
Illinois, United States  
Massachusetts, United States  
Minnesota, United States  
Nevada, United States  
Oklahoma, United States  
Oregon, United States  
Pennsylvania, United States  
Tennessee, United States  
Texas, United States  
Virginia, United States  
China  
Hungary  
Italy  
Poland  
Spain  
Ukraine  
United Kingdom

**IO -centered** No

**Prior IO Allowed** No

**CRC-directed** Yes

**Status** Active, not recruiting

**Drugs** Placebo, Trilaciclib

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**NCT ID** NCT04793958

**Title** Phase 3 Study of MRTX849 With Cetuximab vs Chemotherapy in Patients With Advanced Colorectal Cancer With KRAS G12C Mutation (KRYSTAL-10)

**Phase** Phase 3

**Date Added** 2021-03-11

<b>Location</b>	Alabama, United States Arizona, United States Arkansas, United States California, United States Colorado, United States Connecticut, United States Florida, United States Georgia, United States Illinois, United States Iowa, United States Louisiana, United States Maryland, United States Massachusetts, United States Michigan, United States Minnesota, United States Missouri, United States Nebraska, United States Nevada, United States New Jersey, United States New York, United States North Carolina, United States Ohio, United States Oklahoma, United States South Carolina, United States Tennessee, United States Texas, United States Utah, United States Virginia, United States Washington, United States West Virginia, United States Wisconsin, United States Argentina Australia Austria Belgium Brazil Canada China Colombia Czechia Denmark Finland France Germany Greece Hong Kong Ireland Italy Korea, Republic of Malaysia Mexico Netherlands Poland Portugal Puerto Rico Singapore Spain Taiwan Thailand United Kingdom
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	Yes
<b>CRC-directed</b>	Yes
<b>Status</b>	Recruiting
<b>Drugs</b>	cetuximab, FOLFIRI Regimen, mFOLFOX6 Regimen, MRTX849
<b>NCT ID</b>	NCT04776148
<b>Title</b>	Study of Lenvatinib (MK-7902/E7080) in Combination With Pembrolizumab (MK-3475) Versus Standard of Care in Participants With Metastatic Colorectal Cancer (MK-7902-017/E7080-G000-325/LEAP-017)

<b>Phase</b>	Phase 3
<b>Date Added</b>	2021-03-01
<b>Location</b>	California, United States Georgia, United States Illinois, United States Maryland, United States Michigan, United States Montana, United States Oregon, United States Pennsylvania, United States Virginia, United States Washington, United States Argentina Australia Canada China Denmark Germany Israel Japan Korea, Republic of Russian Federation Spain Taiwan Turkey United Kingdom
<b>IO -centered</b>	No
<b>Prior IO Allowed</b>	No
<b>CRC-directed</b>	Yes
<b>Status</b>	Active, not recruiting
<b>Drugs</b>	Lenvatinib, Pembrolizumab, Regorafenib, TAS-102 (trifluridine and tipiracil)